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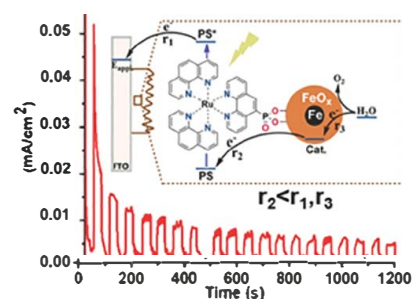
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# Covalent Grafting of Ruthenium Complexes on Iron Oxide Nanoparticles: Hybrid Materials for Photocatalytic Water Oxidation

Quyên T. Nguyen, Elodie Rousset, Van T. H. Nguyen, Vincent Colliere, Pierre Lecante, Wantana Klysubun, Karine Philippot, Jérôme Esvan, Marc Respaud, Gilles Lemerrier,\* Phong D. Tran,\* and Catherine Amiens\*

**ABSTRACT:** The present environmental crisis prompts the search for renewable energy sources such as solar driven production of hydrogen from water. Herein, we report an efficient hybrid photocatalyst for water oxidation, consisting of a ruthenium polypyridyl complex covalently grafted on core/shell Fe@FeO<sub>x</sub> nanoparticles *via* a phosphonic acid group. The photoelectrochemical measurements were performed under 1 sun illumination in 1 M KOH. The photocurrent density of this hybrid photoanode reached 20 μA/cm<sup>2</sup> (applied potential of +1.0 V *vs* reversible hydrogen electrode), corresponding to a turnover frequency of 0.02 s<sup>-1</sup>. This performance represents a 9 fold enhancement of that achieved with a mixture of Fe@FeO<sub>x</sub> nanoparticles and a linker free ruthenium polypyridyl photosensitizer. This increase in performance could be attributed to a more efficient electron transfer between the ruthenium photosensitizer and the Fe@FeO<sub>x</sub> catalyst as a consequence of the covalent link between these two species through the phosphonate pendant group.

**KEYWORDS:** covalent grafting, hybrid photoanode, iron oxide nanoparticles, ruthenium based photosensitizer, water oxidation catalysis



## 1. INTRODUCTION

The 21st century has seen a boom in the development of renewable energy sources, as the world attempts to transition from the serious environmental and health issues associated with the use of non renewable sources such as coal or oil. In this context, hydrogen gas evolved from solar water splitting has attracted tremendous interest as a green fuel.<sup>1</sup> Solar driven water splitting is a thermodynamically uphill process comprising two sluggish half reactions: the oxygen evolution reaction (OER) and the hydrogen evolution reaction (HER). It can be accomplished in a photoelectrochemical cell comprising a photoanode where the OER takes place with solar energy input, and a photocathode (or a simple cathode) where the HER occurs. As the OER is more challenging to achieve, a large research effort focuses nowadays on the development of appropriate photoanodes. In general, a photoanode is constituted of a *photosensitizer* (PS) for photon harvesting and a *catalyst* for accelerating the OER. N type semiconductors such as TiO<sub>2</sub>,<sup>2</sup> BiVO<sub>4</sub>,<sup>3</sup> and α-Fe<sub>2</sub>O<sub>3</sub>,<sup>4</sup> or molecular dyes such as ruthenium tris bipyridine complexes,<sup>5,6</sup> perylene,<sup>7,8</sup> polyheptazine,<sup>9,10</sup> metal free porphyrins,<sup>11</sup> and π conjugated naphthalene benzimidazole polymers<sup>12</sup> have been assayed as PSs in this context. Regarding the OER catalyst, oxides of transition metals such as IrO<sub>2</sub>,<sup>13,14</sup> CoO<sub>x</sub>,<sup>15,16</sup> Co-Pi,<sup>17-19</sup> or FeOOH<sup>20-23</sup> have been investigated, as well as some molecular complexes.<sup>24,25</sup>

To build the photoanode, a dyad comprising a molecular complex catalyst linked to a molecular PS can be deposited on

a transparent electrode.<sup>26,27</sup> The molecular catalyst can also be combined with an N type semi conducting material as a PS. This can be done either by simple physical adsorption or by covalent grafting. This second strategy may lead to photoanodes with improved performance<sup>28</sup> if it can provide (i) an improved electron/hole transfer between the two components and, beyond, an enhanced overall photocatalytic activity and (ii) a more robust system preventing the leaching out of the catalyst.

When a metal oxide catalyst is employed, it can also be combined with an N type semi conducting material acting as a PS. In this case, the two materials are generally interfaced without any specific linker to create a solid–solid junction.<sup>16,19,29-31</sup> Alternatively, a hybrid photoanode can be engineered by covalently grafting a molecular PS onto the surface of the metal oxide catalyst. Similarly, when a molecular catalyst is grafted on a solid PS, such covalent grafting between the molecular PS and solid catalyst is believed to offer (i) an efficient electron/hole transfer between the two components and (ii) a more robust system preventing the leaching out of

the PS. To the best of our knowledge, the first example of such a photoanode for solar water splitting was reported by Mallouk and co workers in 2009, wherein a ruthenium tris bipyridine complex, acting as a PS, was covalently grafted through a malonate linker on  $\text{IrO}_2$  nanoparticles (NPs), used here as a catalyst.<sup>32</sup> Since then, only a few other systems were published.<sup>7,33</sup> Recently, our group also investigated this approach and reported the covalent grafting of a Ru complex PS onto  $\text{Co}_3\text{O}_4$  NPs, using pendant phosphonate linkers and its positive impact on the photoanode efficiency.<sup>34</sup> We then expanded our work toward more environment friendly catalysts, such as iron based systems, considering that iron oxides<sup>35,36</sup> and iron oxyhydroxides<sup>20–23</sup> have been reported to be active and stable electrocatalysts for the OER, but that none of these materials have been used so far to build a hybrid photoanode by combination with a molecular PS.

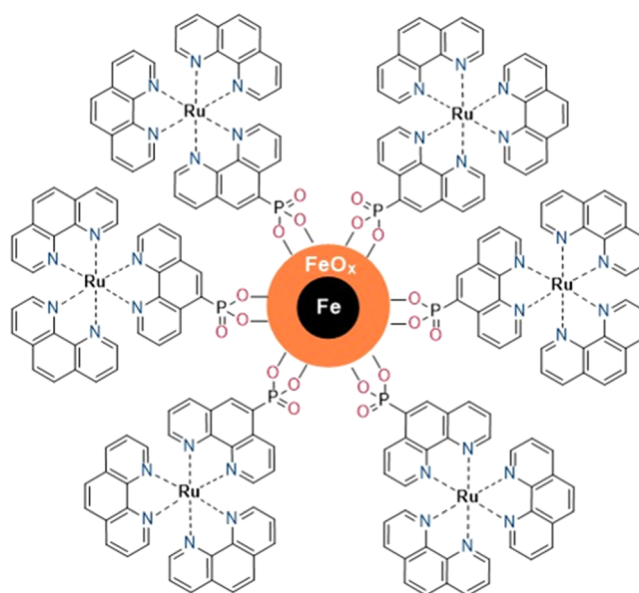
Ruthenium trispolypyridyl complexes are archetypal molecular PSs<sup>37–39</sup> as they present large extinction coefficients over a broad range of the visible spectrum, a long lived excited state, and high luminescent quantum yields. This allows an efficient energy/electron transfer to the catalyst. Their scaffold is also highly tunable, allowing the fine tuning of the optical properties and redox potentials. In comparison with the archetypal  $[\text{Ru}(\text{bpy})_3]^{2+}$  that is mainly used in the development of photo electrocatalytic systems, the phenanthroline based analogues present a wide interest due to their more rigid and conjugated ligands.<sup>40,41</sup> This type of complex is also air stable and displays a high thermal stability allied to a remarkable chemical inertness.<sup>42,43</sup> In our study, a pendant phosphonate group was introduced on the fifth position of one of the 1,10 phenanthroline ligands, as such functional groups are reported to afford a covalent grafting onto iron oxide surfaces.<sup>44–47</sup> This position is usually substituted as it offers efficient charge transfers in photophysical applications, and it is the less sterically hindered in the resulting complex, potentially allowing an easier grafting on the NPs.

We report herein an unprecedented robust hybrid photo catalyst for OER where ruthenium trisphenanthroline complexes (as PSs) are grafted on core/shell iron@iron oxide NPs ( $\text{Fe@FeO}_x$  NPs) used as a catalyst (Figure 1). This hybrid photocatalyst was deposited onto a fluoride doped tin oxide (FTO) electrode to generate a novel hybrid photoanode. Its photocatalytic activity was assayed to evaluate the interest of the covalent coupling between the PS and catalyst.

## 2. RESULTS AND DISCUSSION

### 2.1. Synthesis and Characterization of the OER Catalyst.

In previous studies, we demonstrated that the grafting of a phosphonic group was more efficient on NPs showing a substantial iron oxide layer at their surface.<sup>48</sup> Thus, in this work, we prepared  $\text{Fe@FeO}_x$  core/shell NPs presenting a crystalline iron oxide shell. A two step procedure was followed (Scheme 1). First, body centered cubic (bcc) iron NPs (Fe NPs) were prepared following the procedure reported by Gharbi *et al.*<sup>48</sup> Then, based on the work of Peng *et al.*,<sup>49</sup> the controlled oxidation of these native NPs was performed to generate an iron oxide shell. Analysis by transmission electron microscopy (TEM) showed that the obtained NPs displayed a mean size of  $11.5 \pm 2.3$  nm and a “flower like” morphology (Supporting Information, Figure S1). Their study under high resolution conditions confirmed that this discontinuous shell of lighter contrast (Figure 2a) consisted in well defined iron oxide



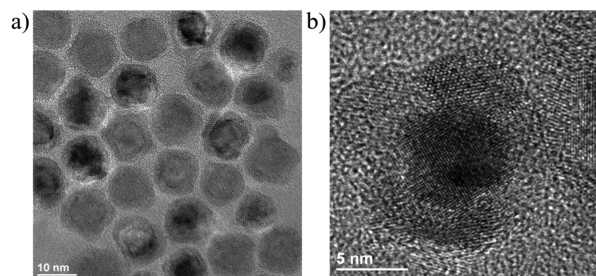
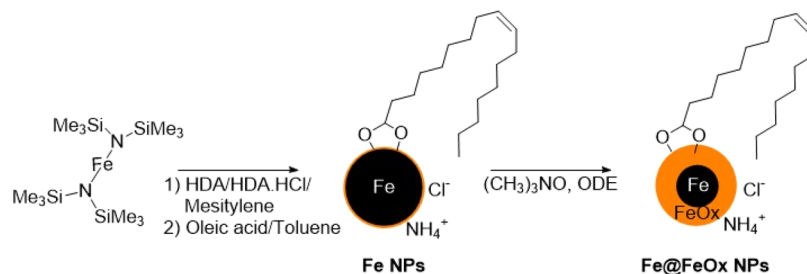
**Figure 1.** Schematic view of the hybrid nanomaterial where Ru(II) tris(1,10 phenanthroline) complexes are grafted onto the  $\text{Fe@FeO}_x$  NPs via a phosphonate group.

crystalline grains (Figure 2b and Supporting Information, Figure S1c,d).

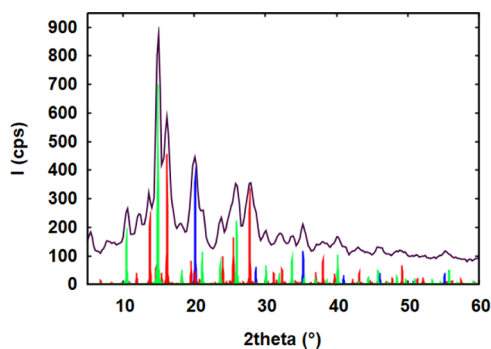
The crystallinity of the oxide shell was further evidenced, on a more statistical basis, through wide angle X ray scattering (WAXS) measurements. The diagram obtained for the  $\text{Fe@FeO}_x$  NPs is presented in Figure 3, together with bcc Fe ( $\alpha$  Fe),  $\gamma$   $\text{Fe}_2\text{O}_3$  (maghemite), and  $\text{NH}_4\text{Cl}$  reference diagrams. Diffraction peaks at 10, 15, and  $26^\circ$  were assigned to  $\text{NH}_4\text{Cl}$ , a product issued from a side reaction between HDA-HCl and hexamethyldisilazane in the first step of the synthesis, as already reported by our group.<sup>50</sup> We also observed peaks assignable to  $\gamma$   $\text{Fe}_2\text{O}_3$  and to bcc Fe, clearly demonstrating the formation of a crystalline oxide shell around the iron core.

The persistence of the iron core was also probed by extended X ray absorption fine structure (EXAFS) at the Fe K absorption edge. Contrarily to WAXS, which places a strong emphasis on large crystalline domains, EXAFS is strictly limited to short range order. On the other hand, this limitation makes it much more sensitive for the analysis of small domains, and contributions not involving Fe atoms would be rejected in principle. A pure Fe foil was measured under the same conditions as  $\text{Fe@FeO}_x$  NPs and additional reference data from the pure  $\gamma$   $\text{Fe}_2\text{O}_3$  were imported from the Farrel Lytle database. Comparing the  $\chi$  function of the  $\text{Fe@FeO}_x$  NPs with those of the two references (Supporting Information, Figure S2), the pattern of  $\gamma$   $\text{Fe}_2\text{O}_3$  was clearly observed with, however, some discrepancies (in the  $8.2\text{--}9.5 \text{ \AA}^{-1}$  range). Interestingly, these discrepancies could be clearly attributed to metallic Fe features. A model including the short Fe–O bonding distance and the longer Fe–Fe distance with an O bridge typical of an iron oxide, as well as the metallic bonding Fe–Fe distance, allowed a good fit of the experimental data, especially in the  $6\text{--}10 \text{ \AA}^{-1}$  range (Table S1, Figure S3). Good agreement was also observed for the Fourier transform (Figures 4 and S4). A model considering more shells could likely provide even better agreement but without providing more insights on this composite sample which likely includes Fe(III) oxide at different degrees of crystallization. For the same reason,

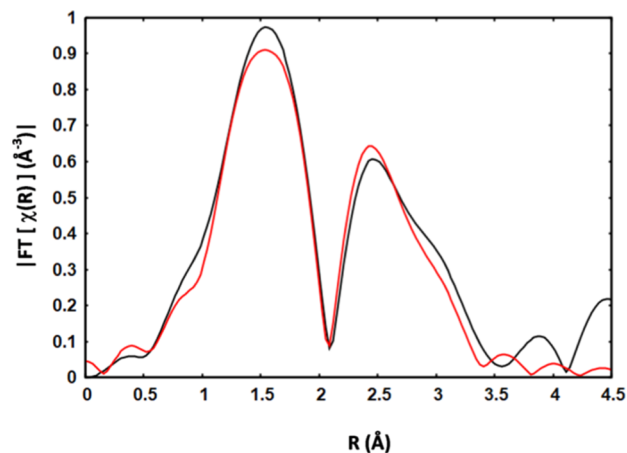
## Scheme 1. Synthetic Pathway of the Fe@FeO<sub>x</sub> Core/Shell NPs



**Figure 2.** Typical HR TEM images of the Fe@FeO<sub>x</sub> NPs with different scale bars: (a) 10 and (b) 5 nm.



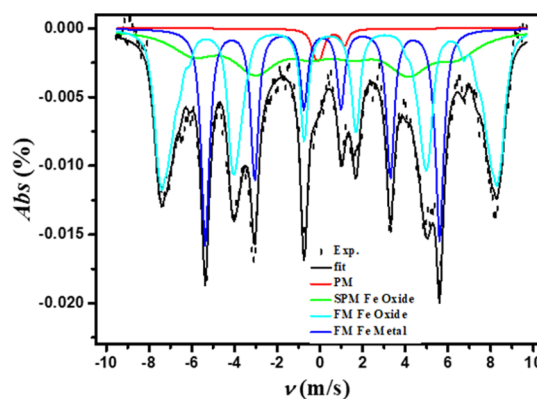
**Figure 3.** WAXS diagram of the Fe@FeO<sub>x</sub> NPs (black solid line) in comparison with those of bcc Fe (PDF 01 071 4648, in blue),  $\gamma$  Fe<sub>2</sub>O<sub>3</sub> (PDF 01 089 5894, in red), and NH<sub>4</sub>Cl (PDF 96 901 0007, in green) references.



**Figure 4.** Magnitude of the Fourier transform of function  $\chi(R)$  from the Fe@FeO<sub>x</sub> NPs (black line) and the best fit achieved (red line).

quantitative evaluation of the Fe/Fe oxide ratio was not attempted even if a mixed organization was demonstrated.

From the Fe 2p spectrum obtained by X ray photoelectron spectroscopy (XPS) (Supporting Information, Figure S5), only binding energies corresponding to Fe(0) (706.7 eV) and Fe(III) in an iron oxide environment (710.3 eV) were observed. This confirmed the presence of a residual Fe core and indicated that the oxide shell was most likely  $\alpha$  or  $\gamma$  Fe<sub>2</sub>O<sub>3</sub>, even if the presence of FeOOH species on the top surface could not be excluded.



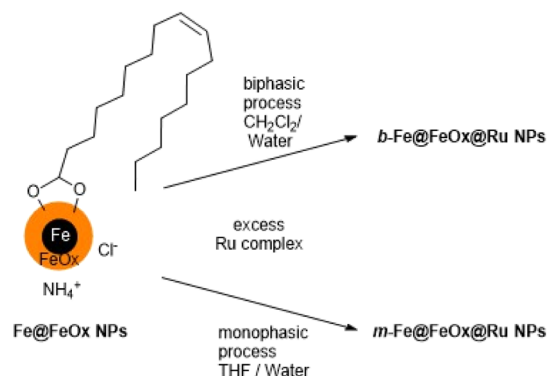
**Figure 5.** Mössbauer spectrum of the Fe@FeO<sub>x</sub> NPs recorded at 80 K (experimental data in black markers), and the best fit obtained (solid line) from a combination of iron and iron oxide contributions (see legend in the figure).

A quantitative evaluation of the Fe/FeO<sub>x</sub> ratio in the NPs and identification of the oxide formed were obtained from the Mössbauer spectrum (Figure 5, Table S2) recorded at 80 K. At this temperature, and given the size of the NPs, relaxation phenomena on the time scale of Mössbauer spectroscopy ( $10^{-11}$  to  $10^{-9}$  s) were initially expected to be negligible. Given the morphology of the NPs, the defined lines observed in the spectrum could be attributed to the contributions of ferromagnetic iron (FM Fe,  $28.9 \pm 0.5\%$ ) and iron oxide phases (FM Fe oxide,  $42.9 \pm 0.5\%$ ). However, to precisely adjust the experimental curve, two other contributions had to be introduced: the first one corresponding to iron oxide in superparamagnetic regime (SPM Fe oxide,  $26.2 \pm 0.5\%$ ) and the second one corresponding to a trace of paramagnetic iron (PM Fe,  $2.0 \pm 0.2\%$ ). Table S2 and Figure S6 display the hyperfine fields, quadrupolar splitting, and the corresponding isomer shifts used to fit the spectrum. The parameters used to fit the iron oxide contributions are in good agreement with those expected for  $\gamma$  Fe<sub>2</sub>O<sub>3</sub> at the nanoscale.<sup>51</sup> The two contributions observed (one FM and the other SPM) could reflect the heterogeneity of the iron oxide shell thickness and polycrystallinity. The Fe(0)/Fe(III) atomic ratio extracted from fitting the Mössbauer spectrum indicated an average shell

thickness of 2.6 nm (Supporting Information, Section S1.5). Thus, the  $\text{Fe@FeO}_x$  NPs can be best described as core/shell  $\text{Fe@}\gamma\text{-Fe}_2\text{O}_3$  NPs with a mean size of  $11.5 \pm 2.3$  nm and an average shell thickness of 2.6 nm. We note that having the  $\gamma\text{-Fe}_2\text{O}_3$  phase on the NP shell is highly desirable as this phase is more catalytically active for the water oxidation reaction than the  $\alpha\text{-Fe}_2\text{O}_3$  phase.<sup>36</sup> We then treated these core/shell  $\text{Fe@}\gamma\text{-Fe}_2\text{O}_3$  NPs with the  $[\text{Ru-PO}(\text{OH})_2]_2\text{Cl}_2$  complex in order to create the hybrid photocatalyst.

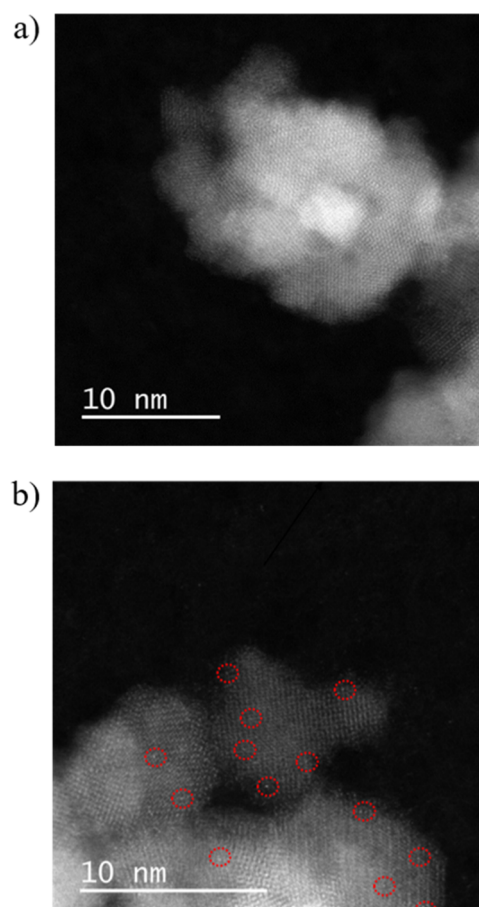
**2.2. Synthesis and Characterization of the Hybrid Photocatalyst.** The synthesis of the 5-phosphonic 1,10-phenanthroline ligand and Ru complex (hereafter referred to as  $[\text{Ru-PO}(\text{OH})_2]_2\text{Cl}_2$ ) were adapted from those reported for the synthesis of 2,2'-bipyridine<sup>52</sup> and of bis-heteroleptic ruthenium(II) complexes<sup>53,54</sup> (see Scheme S1, Figures S8–S11 in the Section S2 of Supporting Information for details). The resulting complex  $[\text{Ru-PO}(\text{OH})_2]_2\text{Cl}_2$  absorbs in the visible domain and is emissive through its <sup>3</sup>MLCT excited state (Supporting Information, Figures S12–S14), showing similar photophysical properties to the  $[\text{Ru}(\text{phen})_3]^{2+}$  reference complex.

**Scheme 2. Synthetic Pathways for the Hybrid Photocatalysts Prepared in a Biphasic Medium (*b*  $\text{Fe@FeO}_x\text{@Ru}$  NPs) or in a Monophasic Medium (*m*  $\text{Fe@FeO}_x\text{@Ru}$  NPs)**



Two synthetic pathways were developed to graft the  $[\text{Ru-PO}(\text{OH})_2]_2\text{Cl}_2$  complex on the  $\text{Fe@FeO}_x$  NPs (Scheme 2). At first, a biphasic method previously used to graft 3-amino-phosphonic acid on similar NPs was followed (Figure S15).<sup>48</sup> However, it afforded only a poor grafting density of the Ru complex on the iron oxide surface (9 Ru complexes per NP, sample hereafter called **b**  $\text{Fe@FeO}_x\text{@Ru}$  NPs, **b** standing for biphasic) (Supporting Information, Section S3.3, Figure S16).

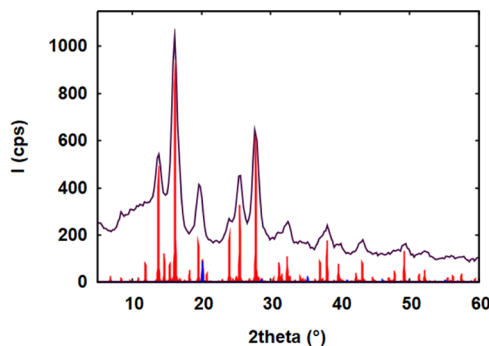
A monophasic method was then implemented, in which  $\text{Fe@FeO}_x$  NPs and  $[\text{Ru-PO}(\text{OH})_2]_2\text{Cl}_2$  were added to a homogeneous THF–water phase (Supporting Information Section S3.2, Figure S17). The magnetic component was recovered from the resulting suspension by magnetic separation. The resulting samples, hereafter referred to as *m*  $\text{Fe@FeO}_x\text{@Ru}$  NPs (*m* standing for monophasic), were washed 5 times with Milli Q water, 1 time with ethanol, and 1 time with diethyl ether to remove the non-grafted ruthenium complex. Analysis of high-resolution high-angle annular dark-field scanning TEM (HR HAADF STEM) images of the *m*  $\text{Fe@FeO}_x\text{@Ru}$  NPs showed that the core/shell morphology of the parent NPs was preserved, but suggested a thicker iron oxide shell (Figure 6). Accordingly, the crystalline iron oxide component was now the dominant feature of the WAXS



**Figure 6.** HR HAADF STEM analysis of the *m*  $\text{Fe@FeO}_x\text{@Ru}$  NPs: (a) typical NP image and (b) atomic resolution micrograph showing the presence of dispersed Ru atoms (highlighted by red circles).

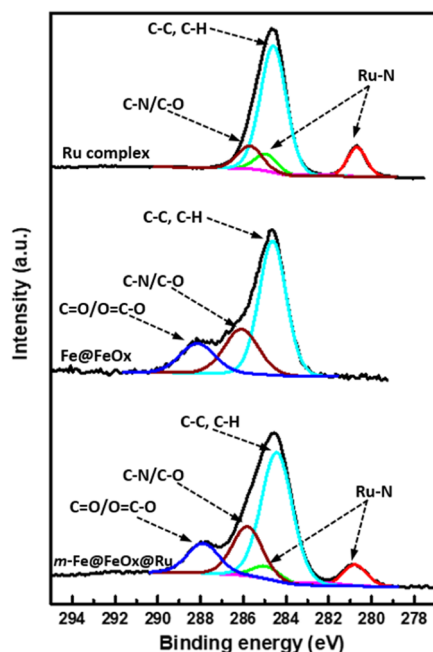
diagram (Figure 7).<sup>55</sup> The width of the peak at  $2\theta = 20^\circ$  and the presence of a peak at  $2\theta = 35^\circ$  suggested a very small contribution of metallic Fe even though it could not be confirmed, nor disproved. The growth of the oxide layer at the expense of the iron core was further confirmed by XPS as in the Fe 2p region only one peak at 710.2 eV, attributed to  $\text{Fe}^{3+}$ , was observed (Supporting Information, Figure S19).

The grafting of the Ru complex on the NPs was first indicated by the presence of the dispersed bright dots in the HR HAADF STEM images of the sample (Figure 6b). As the atomic number *Z* of a Ru atom is higher than that of an Fe



**Figure 7.** WAXS diagram of the *m*  $\text{Fe@FeO}_x\text{@Ru}$  NP sample (black solid line) in comparison with  $\gamma\text{-Fe}_2\text{O}_3$  (PDF 01 089 5894, in red) and bcc Fe (PDF 01071 4648, in blue) references.

atom, these bright dots were tentatively attributed to Ru atoms from the grafted Ru complexes. The integrity of the Ru complexes was evidenced by XPS. Spectra collected in the Ru 3d and C 1s energy ranges on the *m* Fe@FeO<sub>x</sub>@Ru NPs, Fe@FeO<sub>x</sub> NPs, and [Ru-PO(OH)<sub>2</sub>]Cl<sub>2</sub> complex are shown in Figure 8. The free Ru complex showed a Ru 3d<sub>5/2</sub> peak at



**Figure 8.** XPS spectra of the free ruthenium complex [Ru-PO(OH)<sub>2</sub>]Cl<sub>2</sub> (top), Fe@FeO<sub>x</sub> NPs (middle), and *m* Fe@FeO<sub>x</sub>@Ru NPs (bottom) in the C 1s-Ru 3d region (red and green curves: Ru-N 3d<sub>5/2</sub> and 3d<sub>3/2</sub> signals, respectively; cyan curve: C-C/C-H signal; brown curve: C-N/C-O signal; and blue curve: C=O/O=C-O signal).

binding energy of 280.7 eV, being characteristic of a Ru-N environment.<sup>56</sup> The Ru 3d<sub>3/2</sub> peak associated was observed at 285 eV (Supporting Information, Table S3). Two similar peaks were observed for the *m* Fe@FeO<sub>x</sub>@Ru NPs but not for the Fe@FeO<sub>x</sub> NPs. In the *m* Fe@FeO<sub>x</sub>@Ru NPs spectrum, a N 1s peak attributed to the N-Ru bond was observed at 399.8 eV (Supporting Information, Figure S20). Interestingly, we did not observe any contribution from N in an ammonium environment (expected at 401.9 eV<sup>57</sup>) confirming again the elimination of NH<sub>4</sub>Cl during the grafting process, as observed in the WAXS measurements (Supporting Information, Figure S18). Furthermore, in the *m* Fe@FeO<sub>x</sub>@Ru NPs spectrum, the C 1s peak at 284.7 eV could be attributed to C-C/C-H bonds from the phenanthroline ligand and/or from the oleic acid used during the NP synthesis and retained on their surface, whereas the peak at a higher binding energy (285.8 eV) could be attributed to the C-N bond of the phenanthroline ligand and/or the C-O bond of the oleic acid. The persistence of some oleic acid at the surface of the NPs was further confirmed by the peak at 288 eV, assigned to C=O/O=C-O bonds, which was observed in the spectra of both the Fe@FeO<sub>x</sub> NPs and *m* Fe@FeO<sub>x</sub>@Ru NPs (Figure 8 and Supporting Information, Table S3), but not in the spectrum of the Ru complex. The persistence of oleic acid evidences that the exchange of the carboxylic acid coating of the NPs by

phosphonates is difficult at room temperature, even when an excess of phosphonate is introduced.<sup>45</sup>

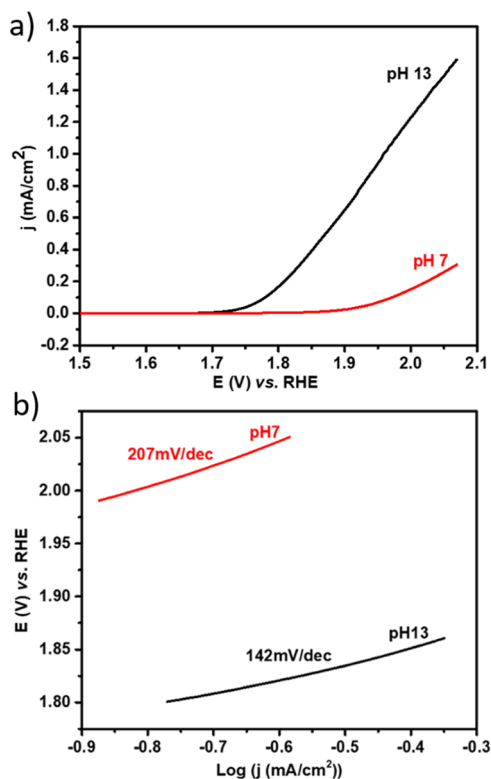
The Ru complex content in the *m* Fe@FeO<sub>x</sub>@Ru NPs was quantitatively determined by inductively coupled plasma optical emission spectroscopy (ICP OES) at ca. 56 Ru complexes per NP against ca. 9 Ru complexes per NP in the *b* Fe@FeO<sub>x</sub>@Ru NPs (see Supporting Information Section S3.3 for calculation details).

All these data clearly provide evidence of the presence of the Ru complex on the surface of the *m* Fe@FeO<sub>x</sub>@Ru NPs although much less Ru complex was grafted *via* the biphasic method than *via* the monophasic one. The higher grafting density obtained by the monophasic method may arise from more efficient interactions between the ruthenium complexes and the NPs due to the miscibility of the solvents, while in the biphasic method such interactions can happen only at the water/dichloromethane interface.

**2.3. Electrocatalytic Activity of Fe@FeO<sub>x</sub> NPs in Water Oxidation.** In the prepared (*m/b*) Fe@FeO<sub>x</sub>@Ru hybrid systems, the role of the molecular complex is to harvest solar energy, whereas the Fe@FeO<sub>x</sub> core aims to accelerate the water oxidation reaction. Thus, before investigating the photocatalytic activity of these hybrid systems, the electrocatalytic activity of the Fe@FeO<sub>x</sub> NPs was evaluated. To this end, a catalyst ink made of Fe@FeO<sub>x</sub> NPs in a EtOH/water solvent mixture together with 1 μL Nafion 5% (wt %) as a linker was drop casted on a FTO electrode at a mass loading density of 1.8 × 10<sup>-4</sup> g/cm<sup>2</sup> (Supporting Information, Figure S21). The resulting catalyst electrode was then assayed in a pH 7 phosphate buffer (KPi 0.1 M) and a pH 13 alkaline electrolyte (NaOH 0.1 M). At pH 13, it operated with an onset potential of ~1.75 V *versus* reversible hydrogen electrode (RHE) and a Tafel slope value of 142 mV/decade could be deduced from the polarization curves (Figure 9, black trace). At pH 7, it showed a much higher onset potential of ~1.95 V as well as a higher Tafel slope value: 207 mV/decade (Figure 9, red trace). This assay demonstrated that the Fe@FeO<sub>x</sub> NPs is an active catalyst for the OER, and that it is significantly more active in an alkaline solution.

**2.4. Photoelectrochemical Water Oxidation Catalysis with Fe@FeO<sub>x</sub>@Ru Hybrid Electrodes.** We investigated the photocatalytic activity of the Fe@FeO<sub>x</sub>@Ru NPs hybrid nanomaterials to understand to what extent a covalent linkage between the molecular PS and the nanocatalyst can benefit its overall operation. To this end, an ink was prepared from the Fe@FeO<sub>x</sub>@Ru NPs samples and deposited onto a FTO electrode as described above for the Fe@FeO<sub>x</sub> NPs. The light source was provided by a Xe arc lamp equipped with a UV filter. We employed the back illumination mode in which the incident light went through the transparent FTO electrode to the catalyst. The incident light intensity reaching the FTO electrode was set to 100 mW/cm<sup>2</sup>.

Figure 10a shows a simplified overview of the electron transfers involved during the operation of the hybrid photocatalyst. In this model, the overall oxygen production rate is determined by the slowest step of the elemental reactions 1, 2, and 3, in which: (i) *r*<sub>1</sub> represents the rate of the oxidation of the excited photosensitizer (PS\*) by an oxidative potential applied to the FTO electrode. Thus, *r*<sub>1</sub> is expected to increase when an increasing oxidative potential is applied; (ii) *r*<sub>2</sub> represents the electron transfer rate from the catalyst to the oxidized photosensitizer (PS<sup>+</sup>) (*i.e.*, the hole transfer rate from PS<sup>+</sup> to the catalyst), and (iii) *r*<sub>3</sub> represents the O<sub>2</sub> evolution



**Figure 9.** (a) Polarization curves of an  $\text{Fe@FeO}_x$  NP electrode recorded in pH 7 (red line) and pH 13 (black line) electrolyte solutions. Potential scan rate: 5 mV/s. (b) Tafel plots of the  $\text{Fe@FeO}_x$  NP electrode at pH 7 (red line) and pH 13 (black line).

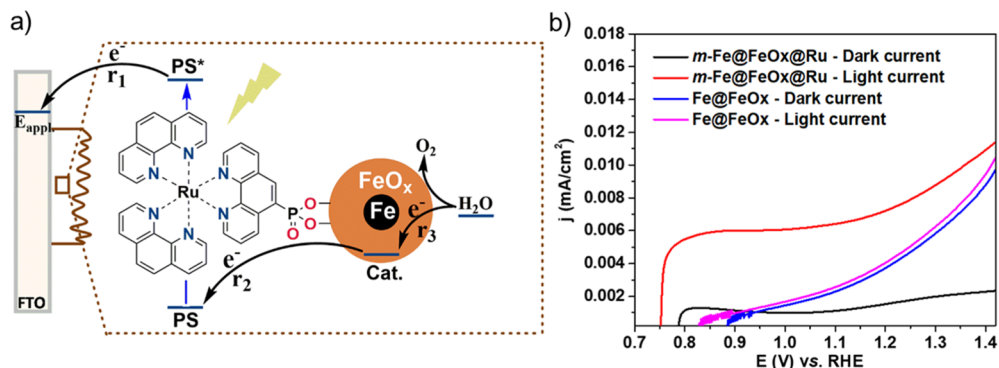
rate on the surface of the water oxidation catalyst, that is, on the surface of the  $\text{Fe@FeO}_x$  component of the hybrid nanomaterial.

Herein, the photocatalytic activity of the  $m \text{Fe@FeO}_x\text{@Ru}$  NP electrode for the water oxidation reaction was assayed at pH 13, where  $r_3$  is assumed to be optimal. In these conditions, an  $\text{Fe@FeO}_x$  control electrode was not photocatalytically active, showing negligible photogenerated current under 1 sun illumination (Figure 10b, blue and pink traces). A negligible current density was also recorded for the  $m \text{Fe@FeO}_x\text{@Ru}$  NP electrode under dark condition at applied potentials up to 1.5 V versus RHE (Figure 10b, black trace). Under light

illumination, the onset potential for generating a measurable photocurrent was found to be 0.75 V versus RHE. An almost constant photocurrent density was observed when increasing the applied potential up to 1.1 V versus RHE. This means that the overall rate of water oxidation remained unchanged in this potential window, even if  $r_1$  should increase when increasing the applied oxidative potential. Therefore, the charge transfer between the Ru PS and the  $\text{Fe@FeO}_x$  nanocatalyst, namely  $r_2$ , limited the whole operation in this potential range.

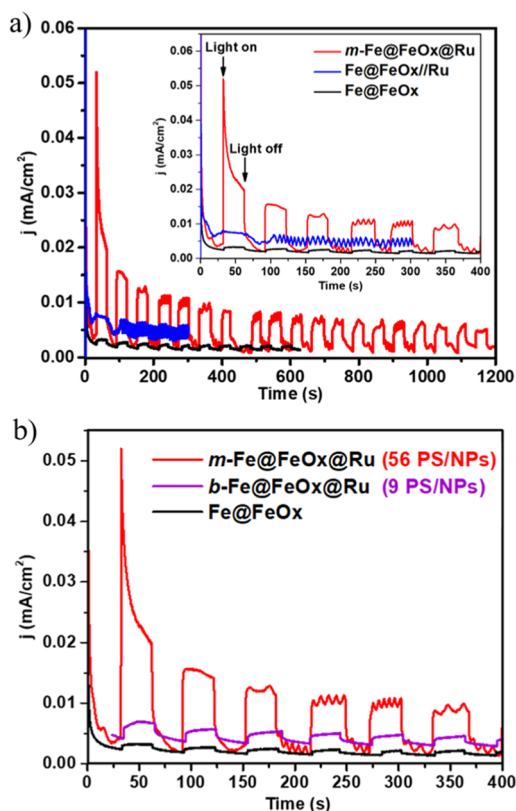
In order to emphasize the benefit of engineering a covalent grafting between the PS and the catalyst, we also recorded the photocatalytic activity of a simple mixture of  $\text{Fe@FeO}_x$  NPs and ruthenium tris phenanthroline chloride,  $[\text{Ru}(\text{phen})_3]\text{Cl}_2$ , in the same Ru per NP ratio (i.e., 56) as in the  $m \text{Fe@FeO}_x\text{@Ru}$  NP sample, hereafter labeled  $\text{Fe@FeO}_x\text{/Ru}$ .  $[\text{Ru}(\text{phen})_3]\text{Cl}_2$  was chosen as a reference PS given its comparable photophysical properties to those of  $[\text{Ru}-\text{PO}(\text{OH})_2]\text{Cl}_2$  (see Supporting Information, Figure S13) and was expected to adsorb only physically onto the  $\text{Fe@FeO}_x$  NPs given the absence of pendant grafting groups. A control electrode was also fabricated by loading only  $\text{Fe@FeO}_x$  NPs at an identical mass loading density without adding any Ru complex. Figure 11a shows the  $I-t$  curves recorded at 1.0 V versus RHE under chopped light mode using electrodes made of  $m \text{Fe@FeO}_x\text{@Ru}$  NPs (red trace),  $\text{Fe@FeO}_x\text{/Ru}$  (blue trace), and  $\text{Fe@FeO}_x$  NPs (black trace). The  $\text{Fe@FeO}_x$  NP photoanode generated a negligible photocurrent density of ca.  $0.5 \mu\text{A}/\text{cm}^2$ .  $\text{Fe@FeO}_x\text{/Ru}$  (simple mixture) showed a higher photocurrent density ( $2.3 \mu\text{A}/\text{cm}^2$ ) demonstrating the improvement in light harvesting brought by the Ru complex. Interestingly, a significantly higher photocurrent density of ca.  $20 \mu\text{A}/\text{cm}^2$  was achieved for the  $m \text{Fe@FeO}_x\text{@Ru}$  NP electrode (where the Ru complex is covalently grafted on the  $\text{Fe@FeO}_x$  nanocatalyst). This value corresponds to ca. 9 fold and 40 fold higher activities than those obtained with the  $\text{Fe@FeO}_x\text{/Ru}$  and  $\text{Fe@FeO}_x$  electrodes, respectively. This result clearly demonstrates, as suggested in the literature,<sup>34,58</sup> the usefulness of a covalent grafting between the Ru PS and the  $\text{Fe@FeO}_x$  catalyst to promote the charge transfer and therefore increase the overall photocatalytic activity.

The robustness of our hybrid system was evaluated by repeating the on-off light illumination cycle. The  $m \text{Fe@FeO}_x\text{@Ru}$  NP photoanode was found to still produce ~50% of the initial photocurrent density (determined in the first on-off



**Figure 10.** (a) Simplified scheme for the photocatalytic water oxidation where PS and PS\* stand for the photosensitizer in its ground and excited states, respectively, and cat stands for the catalyst (here the  $\text{Fe@FeO}_x$  NPs). (b)  $I-V$  curves recorded on the  $m \text{Fe@FeO}_x\text{@Ru}$  NP (dark: black trace; illumination: red trace) and the  $\text{Fe@FeO}_x$  NP (dark: blue trace; illumination: pink trace) photoanodes immersed in a pH 13 (NaOH 0.1 M) electrolyte. Potential scan rate: 10 mV/s, 1 sun illumination.





**Figure 11.** (a)  $I-t$  curves recorded on  $m$   $\text{Fe@FeO}_x\text{@Ru}$  NP (red trace),  $\text{Fe@FeO}_x\text{//Ru}$  (blue trace), and  $\text{Fe@FeO}_x$  NP (black trace) photoanodes; (b)  $I-t$  curves recorded on  $m$   $\text{Fe@FeO}_x\text{@Ru}$  NP (red trace),  $b$   $\text{Fe@FeO}_x\text{@Ru}$  NP (purple trace), and  $\text{Fe@FeO}_x$  NP (black trace) photoanodes. Applied potential: 1.0 V *vs* RHE. Electrolyte: pH 13 NaOH solution. Light incident power intensity: 100  $\text{mW}/\text{cm}^2$ .

cycle) after 20 min assay, whereas the non bonded  $\text{Fe@FeO}_x\text{//Ru}$  counterpart did not generate any noticeable photocurrent from the second on–off cycle onward. Thus, the covalent grafting significantly improves the stability of the  $m$   $\text{Fe@FeO}_x\text{@Ru}$  NP hybrid system. The degradation of the non bonded  $\text{Fe@FeO}_x\text{//Ru}$  system can be attributed to a fast detachment of  $[\text{Ru}(\text{phen})_3]\text{Cl}_2$  from the electrode surface because of the lack of covalent bonding and the high solubility of this complex in water. It can be also attributed to a rapid decomposition of  $[\text{Ru}(\text{phen})_3]\text{Cl}_2$  due to the lack of an efficient electron communication with the  $\text{Fe@FeO}_x$  NP.

As described above, the  $b$   $\text{Fe@FeO}_x\text{@Ru}$  NPs had a much lower Ru complex grafting density than the  $m$   $\text{Fe@FeO}_x\text{@Ru}$  NPs, that was, 9 Ru complexes *versus* 56 Ru complexes per NP. At identical mass loading density, the  $b$   $\text{Fe@FeO}_x\text{@Ru}$  NP photoanode showed a 7 fold lower photocurrent density than the  $m$   $\text{Fe@FeO}_x\text{@Ru}$  NP photoanode did, that was  $\sim 3$  *versus*  $\sim 20$   $\mu\text{A}/\text{cm}^2$  (Figure 11b, purple and red traces). However, comparable photocatalytic performances were deduced for these two photoanodes when normalizing the catalytic activity per number of PS, that was, per number of grafted Ru complex (see Supporting Information, Section S4 for details on normalization), thus confirming that the rate limiting step involved the Ru complex. Turnover frequencies per Ru complex ( $\text{TOF}_{\text{Ru}}$ ) of 0.015 and 0.02  $\text{s}^{-1}$  were calculated for the photoanodes prepared with the  $b$   $\text{Fe@FeO}_x\text{@Ru}$  NPs and  $m$   $\text{Fe@FeO}_x\text{@Ru}$  NPs, respectively. The similarity of these values further confirmed that the charge transfer between the

Ru PS and the  $\text{Fe@FeO}_x$  nanocatalyst was the main limiting factor. A higher density of the Ru PS would thus increase the overall performance of the system.

Looking back at the first on–off light illumination cycle recorded with the  $m$   $\text{Fe@FeO}_x\text{@Ru}$  NPs, a high transient photocurrent ( $\Delta J_{\text{transient}}$ ) of 33  $\mu\text{A}/\text{cm}^2$  (Supporting Information, Figure S22) was observed. This suggested an important charge recombination within the system.<sup>59,60</sup> More studies are required to understand this phenomenon. For now, we tentatively attribute this recombination to the quenching of the (luminescent)  $^3\text{MLCT}$  excited state of the Ru PS<sup>61</sup> by the oxygen molecules which are generated locally on the surface of the  $\text{Fe@FeO}_x$  NPs. Such a quenching has been indeed demonstrated in solution, as mentioned earlier in this study (Supporting Information, Figure S14). If so, quickly removing oxygen from the catalyst surface would reduce the charge recombination and thus represents a good strategy to increase the performance of the hybrid photocatalyst.

### 3. CONCLUSIONS

This study reports the successful construction of an unprecedented hybrid photocatalyst for the solar driven water splitting by covalently grafting a Ru phenanthroline PS onto  $\text{Fe@FeO}_x$  core/shell NPs through a phosphonate linker. Well controlled  $\text{Fe@FeO}_x$  core/shell NPs of  $ca. 11.5 \pm 2.3$  nm in diameter and with a  $\gamma$   $\text{Fe}_2\text{O}_3$  oxide shell of  $ca. 2.6$  nm were synthesized. These NPs showed a better  $\text{O}_2$  evolution electrocatalytic activity in alkaline conditions than in neutral ones, with an onset potential of 1.75 V and a Tafel slope value of 142 mV/decade at pH 13. Two synthetic pathways were investigated to graft the Ru PS at the surface of the  $\text{Fe@FeO}_x$  NPs, *via* monophasic and biphasic processes in THF/ $\text{H}_2\text{O}$  or  $\text{CH}_2\text{Cl}_2/\text{H}_2\text{O}$  solvent mixtures, respectively. The monophasic process was found to be more efficient as it provided a higher grafting density at the surface of the NPs, 7 fold higher than that obtained by the biphasic process (56 and 9 Ru complexes per NP for the monophasic and biphasic processes, respectively). Finally, the efficient photocatalytic activity of the hybrid system prepared in monophasic conditions was demonstrated. The results obtained evidenced that the charge transfer between the grafted Ru PS and the  $\text{Fe@FeO}_x$  nanocatalyst was the bottleneck in the operation of these hybrid systems. At an applied potential of +1.0 V *versus* RHE and under a simulated 1 sun illumination, a catalytic rate of 0.02  $\text{O}_2$   $\text{Ru}^{-1} \text{s}^{-1}$  was observed. Thus, increasing the grafting density of the Ru PS could be a way to improve the overall performance. Moreover, the covalent grafting was found to not only improve the photocatalytic activity (by *ca.* 9 fold compared with a system based on a simple physical adsorption) but also to significantly improve the stability of the photocatalyst. However, the charge recombination within the system remained important as evidenced by a rather high transient current. This may result from the quenching of the excited state of the Ru(II) complex by  $\text{O}_2$  molecules accumulating on the surface of the  $\text{Fe@FeO}_x$  catalyst until a steady state in their evolution is reached. If so, a better evacuation of  $\text{O}_2$  might improve the overall performance of the hybrid photocatalyst as well.

The covalent grafting of the hybrid  $m$   $\text{Fe@FeO}_x\text{@Ru}$  NPs on the FTO electrode is now envisaged to avoid any possible leaching of the catalyst and to limit back electron transfer events happening during photocatalytic conditions.

## 4. EXPERIMENTAL SECTION

**4.1. Materials.** All chemicals used in the synthesis of the ligand and ruthenium(II) complex were purchased from commercial suppliers and used without further purification. 5-Bromo-1,10-phenanthroline<sup>62–64</sup> and *cis* [Ru(phen)<sub>2</sub>Cl<sub>2</sub>]<sup>53,54</sup> were synthesized as previously reported.

[Fe[N(SiMe<sub>3</sub>)<sub>2</sub>]<sub>2</sub>]<sub>2</sub> was purchased from Nanomeps. Hexadecylamine (HDA) (98%), oleic acid (99%), hydrogen chloride solution, 2 N in diethylether, trimethylamine *N* oxide (99%), *tert* butyl nitrite (90%), tetrabutylammonium hexafluorophosphate (98%), and Nafion (5 wt % in a mixture of lower aliphatic alcohols and water) were purchased from Sigma Aldrich and used without any purification. Hexadecylammonium chloride (HDA·HCl) was prepared as previously reported.<sup>65</sup> Mesitylene (99.7%) was purchased from Sigma Aldrich and distilled over sodium under an argon atmosphere. Octadecene (ODE) (90%) was purchased from Sigma Aldrich and dried over activated molecular sieve with a pore size of 0.4 nm under an argon atmosphere. Toluene, CH<sub>2</sub>Cl<sub>2</sub>, and diethyl ether were collected from an MBraun solvent purification system. Ethanol, absolute grade, was purchased from Sigma Aldrich and distilled over magnesium under an argon atmosphere. All the solvents were used after degassing by the freeze–pump–thaw technique. Milli Q water with conductivity at 28.2 Ω was used. FTO slides were cleaned by sonication for 30 min in acetone, 30 min in EtOH and then dried at room temperature for at least 1 h before use.

**4.2. Synthesis of the [Ru(phen)<sub>2</sub>(phen-PO(OH)<sub>2</sub>)]Cl<sub>2</sub> Complex—[Ru–PO(OH)<sub>2</sub>]<sub>2</sub>Cl<sub>2</sub>.** **4.2.1. Synthesis of 5-P(O)(OH)<sub>2</sub>-1,10-Phenanthroline (phen-PO(OH)<sub>2</sub>).** Under an argon atmosphere, 200 mg of the already described 5-bromo-1,10-phenanthroline<sup>63</sup> (0.77 mmol) in toluene, 120 μL of commercially available HP(O)(OEt)<sub>2</sub> (0.93 mmol), 130 μL of triethylamine (0.93 mmol), 13 mg of Pd(OAc)<sub>2</sub> (0.058 mmol, 7%), and 65 mg of the ligand 1,1'-bis(diphenylphosphino)ferrocene (dppf) as the co-catalyst (0.12 mmol) were stirred at reflux overnight. The starting brownish suspension turned to an orange solution. The solvent was evaporated to yield the crude material, which was purified by column chromatography on silica with a CH<sub>2</sub>Cl<sub>2</sub>/MeOH/NH<sub>3</sub>·H<sub>2</sub>O (96/2/2) mixture as an eluent. 55 mg of the desired compound (around 0.17 mmol, yield = 22%) was obtained, already partially deprotected and used as obtained.

**4.2.2. Synthesis of [Ru(phen)<sub>2</sub>(phen-PO(OH)<sub>2</sub>)](Cl)<sub>2</sub>.** Under an argon atmosphere, 20 mg of the ligand phen-PO(OH)<sub>2</sub> (0.057 mmol) and 1 equiv of the precursor complex [Ru(Phen)<sub>2</sub>Cl<sub>2</sub>] (31.5 mg) were dissolved in 3 mL of degassed EtOH. The mixture was heated at reflux overnight. After evaporation of the solvent to dryness, 45 mg of an orange/red powder was obtained (yield = 10%). <sup>1</sup>H NMR (D<sub>2</sub>O, 500 MHz): δ 9.04 (1H, d, *J* = 5 Hz), 8.48 (6H, d, *J* = 5 Hz), 8.10 (4H, s), 8.08–7.95 (6H, m), 7.58–7.45 ppm (6H, m); <sup>31</sup>P NMR (D<sub>2</sub>O, 202 MHz): δ 7.13 ppm (P–OH, d, <sup>2</sup>*J* = 40 Hz); HRMS (ESI): *m/z* calcd for RuC<sub>36</sub>H<sub>24</sub>N<sub>6</sub>O<sub>3</sub>PRu – H<sup>+</sup>, 721.0691 [M – H<sup>+</sup>]; exp., 721.0698. Anal. Calcd for [Ru(Phen)<sub>2</sub>(PhenP(O)(OH)<sub>2</sub>)]Cl<sub>2</sub>·7H<sub>2</sub>O (*M<sub>w</sub>* = 918.7 g/mol): C, 46.9; H, 3.8; and N, 9.2. Found: C, 47.1; H, 4.3; N, 9.2.

**4.3. Synthesis of NPs.** **4.3.1. Synthesis of Fe NPs.** Iron NPs were synthesized following a procedure reported by Gharbi *et al.*<sup>48</sup> with a minor modification. [Fe[N(SiMe<sub>3</sub>)<sub>2</sub>]<sub>2</sub>] (1.9 g, 2.5 mmol), HDA (2.4 g, 10 mmol), and HDA·HCl (2.1 g, 7.5 mmol) were dissolved in 100 mL mesitylene in a Fischer–Porter bottle in a glovebox. The dark brown solution was then immersed in an oil bath pre-heated at 150 °C and kept at this temperature for 65 h. The reaction mixture was then cooled to room temperature and the mesitylene was removed by evaporation under reduced pressure. The iron NPs were dispersed in a solution of 8 mL oleic acid in 100 mL of toluene. The quantity of oleic acid was added in stoichiometric proportion *versus* the amine present in the crude mixture at the end of the reaction. Overnight, the NPs precipitated. The particles were then collected by magnetic separation and washed with toluene (4 × 50 mL), then with ethanol (3 × 50 mL) and afterward dried under vacuum. The NPs were kept inside a glovebox before use (recovered mass: 304 mg, ICP OES: Fe = 82 wt %).

**4.3.2. Synthesis of Fe@FeO<sub>x</sub> NPs.** The iron oxide layer was achieved following the reported protocol<sup>49</sup> with minor modifications. In brief, 0.18 mmol of (CH<sub>3</sub>)<sub>3</sub>NO (14 mg) were dispersed in 20 mL of ODE and flushed with argon for 30 min in a Fischer–Porter bottle. The mixture was heated up to 130 °C for 30 min. In another Fischer–Porter bottle, 160 mg of Fe NPs were dispersed in 4 mL of ODE and then this mixture was transferred to the (CH<sub>3</sub>)<sub>3</sub>NO dispersion using a Teflon canula avoiding air exposure and heated for 2 h at 130 °C. Then, the mixture was heated up to 250 °C for 30 min, cooled to room temperature and opened in air. 25 mL of 1-propanol was added to this mixture and the black powder was collected using a magnet. The powder was washed with diethyl ether (1 × 30 mL) and then dried in air. (Recovered mass: 147 mg, ICP OES: Fe = 86.95 wt %).

**4.3.3. Synthesis of Fe@FeO<sub>x</sub>@Ru Hybrid Nanomaterial by the Biphasic Method—*b*-Fe@FeO<sub>x</sub>@Ru NPs.** 50 mg of Fe@FeO<sub>x</sub> NPs were added to 15 mL of CH<sub>2</sub>Cl<sub>2</sub> in a closed glass tube and sonicated for 5 min. Then, 10 mL of a 2.96 × 10<sup>−4</sup> M solution of [Ru(phen)<sub>2</sub>(phen-PO(OH)<sub>2</sub>)]Cl<sub>2</sub>·7H<sub>2</sub>O in Milli Q water was added. The mixture was mechanically stirred for 3 days till the aqueous phase became completely black, indicating that the transfer was efficient. The aqueous phase was then collected and purified by repeated magnetic precipitation–solvent removing–redispersion steps and washed with Milli Q water (5 × 30 mL), then with ethanol (1 × 30 mL) and diethyl ether (1 × 30 mL). The *b*-Fe@FeO<sub>x</sub>@Ru NP sample was dried in air before characterization. Recovered mass: 39.2 mg.

**4.3.4. Synthesis of Fe@FeO<sub>x</sub>@Ru Hybrid Nanomaterial by the Monophasic Method—*m*-Fe@FeO<sub>x</sub>@Ru NPs.** 21 mg of Fe@FeO<sub>x</sub> NPs were dispersed in 15 mL THF using sonication for 20 min. 10 mL of a 2.28 × 10<sup>−4</sup> M solution of [Ru(phen)<sub>2</sub>(phen-PO(OH)<sub>2</sub>)]Cl<sub>2</sub>·7H<sub>2</sub>O in Milli Q water was added to this suspension. The reaction mixture was mechanically stirred at room temperature for 3 days. The *m*-Fe@FeO<sub>x</sub>@Ru NP sample was purified as described above for the biphasic method. Recovered mass: 17.5 mg.

**4.4. Preparation of Photoanodes.** Catalyst inks were prepared by sonicating 3 mg of each catalyst in 1 mL of a 1/4 (v/v) EtOH/H<sub>2</sub>O solvent mixture together with 1 μL of Nafion 5 wt % as a linker.

*m*-Fe@FeO<sub>x</sub>@Ru NP, *b*-Fe@FeO<sub>x</sub>@Ru NP, and Fe@FeO<sub>x</sub> NP photoanodes were prepared by depositing 17 μL of the respective catalyst ink onto a FTO electrode (*S* = 0.283 cm<sup>2</sup>, loading density of 1.8 × 10<sup>−4</sup> g/cm<sup>2</sup>) followed by a mild annealing process at 100 °C (a temperature at which the Ru complex is stable) in an oven for 5 h before testing their photocatalytic activity.

**4.4.1. Preparation of Non-bonded Fe@FeO<sub>x</sub>/Ru Photoanode for Control Experiment.** A 1/4 (v/v) EtOH/H<sub>2</sub>O solvent mixture together with 1 μL of Nafion 5 wt % was used to prepare a stock solution of [Ru(phen)<sub>3</sub>]Cl<sub>2</sub> with concentration 3.8 × 10<sup>−5</sup> M. A catalyst ink was prepared by the dispersion of Fe@FeO<sub>x</sub> NPs (3 mg) in 1 mL of this stock solution. Then, this ink was deposited onto a FTO electrode with a loading amount of 1.8 × 10<sup>−4</sup> g/cm<sup>2</sup>, followed by a mild annealing process at 100 °C for 5 h (a temperature at which the [Ru(phen)<sub>3</sub>]Cl<sub>2</sub> complex is stable).

**4.5. Photoelectrochemical Study.** The photocatalytic activity was assayed in NaOH electrolyte solution, pH = 13. A conventional three-electrode configuration was used with the photoanode (as prepared above) as a working electrode, Ag/AgCl/1 M KCl as a reference electrode, and a Pt wire as a counter electrode.

Light illumination was provided by a Xe arc lamp (LOT, Germany) with a 420 nm cutoff filter. The incident intensity was set to 100 mW/cm<sup>2</sup>. Back side illumination was conducted, in which the incident light went through the transparent FTO electrode to the deposited catalyst on the photoanode. The potential polarization was conducted with a potential scan rate of 10 mV/s.

Potentials are quoted against the RHE using the following eq 1

$$E_{\text{RHE}} = E_{\text{Ag/AgCl/KCl 3 M}} + 0.059 \times \text{pH} + E_{\text{Ag/AgCl}}^0 \quad (1)$$

where the pH is 13 because the measurements were performed in 0.1 M NaOH,  $E_{\text{Ag/AgCl}}^0$  is 0.21 V *versus* RHE.

The current density (*j*, mA/cm<sup>2</sup>) was calculated by normalization to the geometric surface area of the FTO electrode (*S* = 0.283 cm<sup>2</sup>).

**4.6. Other Characterizations.** The NP samples were characterized by different techniques in conventional (TEM) and high resolution (HR TEM) modes, atom resolved microscopy (ARM), WAXS, EXAFS, XPS, ICP OES, and Mössbauer spectroscopy.

**4.6.1. TEM, HR-TEM, and ARM.** These analyses were performed at the “Centre de Microcaractérisation Raymond Castaing” in Toulouse. The shape, size, and crystallinity of the NPs were examined in conventional TEM (JEOL, JEM 1011 microscope, operating at 100 kV, and a point resolution of 0.45 nm) and high resolution HAADF STEM (JEM, ARM200F microscope, operating at 200 kV, and a point resolution of 0.19 nm). TEM grids were prepared by casting a drop of the NP suspension onto a copper grid (400 hexagonal meshes, carbon coated). Then, the grids were further dried under vacuum for at least one night before introduction into the microscopy chamber. Size distributions were acquired by measuring a minimum of 250 objects using open source ImageJ software. Sizes are given as mean  $\pm$  standard deviation according to a Gaussian fit of the corresponding size distribution. Fast Fourier transform analysis was carried out with Digital Micrograph software.

**4.6.2. X-ray Photoelectron Spectroscopy.** XPS measurements were performed at CIRIMAT ENSIACET in Toulouse on a Thermo Scientific K Alpha spectrometer equipped with a monochromatized Al K $\alpha$  ( $h\nu = 1486.6$  eV) source. The X ray spot size was 400  $\mu\text{m}$ . The pass energy was fixed at 30 eV with a step of 0.1 eV for core levels and 160 eV for surveys (step 1 eV). The spectrometer energy calibration was done using the Au 4f $_{7/2}$  (83.9  $\pm$  0.1 eV) and Ag 3d $_{5/2}$  (368.2  $\pm$  0.1 eV) photoelectron lines. XPS spectra were recorded in direct mode N (Ec) and the background signal was removed using the Shirley method. XPS high resolution spectra were recorded in order to extract the chemical environments of the studied species.

**4.6.3. Inductively Coupled Plasma-Optical Emission Spectroscopy.** Fe and Ru contents were determined by ICP OES using a PerkinElmer Optima 2100 DV instrument at LCC CNRS Toulouse. The samples were first digested into a mixture of HCl/HNO $_3$  (3:1 v/v) and then diluted with Milli Q water.

**4.6.4. Wide-Angle X-ray Scattering.** WAXS measurements were carried out at CEMES CNRS in Toulouse. The samples were sealed in 1.0 mm diameter Lindemann glass capillaries under an Ar atmosphere. The X ray scattering intensity measurements were performed using a dedicated two axis diffractometer equipped with a high energy resolution solid state detector allowing for the removal of the fluorescence from iron at the measurement step by electronic filtering, using the molybdenum K $\alpha$  [0.071069 nm] radiation monochromatized by a flat graphite crystal. Time for data collection was typically 20 h for a set of 457 measurements collected at room temperature in the range  $0^\circ < \theta < 65^\circ$  for equidistant  $s$  values [ $s = 4\pi(\sin \theta/\lambda)$ ]. Radial distribution functions were obtained after Fourier transformation of the corrected and reduced data.

**4.6.5. Extended X-ray Absorption Fine Structure.** EXAFS spectra were recorded at the Fe K edge (7112 eV) at room temperature in a transmission mode on a BL8 beamline at Synchrotron Light Research Institute (SLRI, Nakhon Ratchasima, Thailand), using a Ge(220) double crystal monochromator. All data analysis was performed using ATHENA and ARTEMIS softwares.<sup>60</sup> Reference data from the pure  $\gamma$  Fe $_2$ O $_3$  were imported from the Farrel Lytle database maintained by the International X ray Absorption Society.

**4.6.6. Mössbauer Spectroscopy.** The Mössbauer spectrum was obtained using a spectrometer running in the triangular symmetric mode for the velocity and a radioactive source of 1.85 GBq (50 mCi) Co $^{57}$  diffused into a Rh matrix at LCC CNRS Toulouse. The temperature was regulated at 80 K in an Oxford cryostat. In the figures, dots represent the experimental data and continuous lines represent the curves corresponding to the four contributions used to fit the experimental data. The global fitting curve is also displayed as a solid line. Each contribution is defined by its isomer shift with respect to bcc iron ( $\delta_0$ ), quadrupolar splitting ( $\Delta = \epsilon_0$ ), and a distribution of hyperfine fields ( $\mu_0 H_{\text{hyp}}$ ). We have considered the same width ( $\Gamma$ ) of the Lorentzian lines for all the contributions,  $\Gamma = 0.4$  mm/s. The height ratios of line -1 to line -3  $h_1/h_3$  and line -2 to line -3  $h_2/h_3$  have been fixed for the elemental sextet to the theoretical values

expected for randomly oriented systems, that is,  $h_1/h_3 \approx 3$  and  $h_2/h_3 \approx 2$ .

## ■ ASSOCIATED CONTENT

### ● Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acsami.1c15051>.

Synthesis and characterization of [Ru-PO(OH) $_2$ ]Cl $_2$  complex; complementary data on Fe@FeO $_x$  NPs and Fe@FeO $_x$ @Ru NPs; and complementary data on water oxidation catalysis (PDF)

## ■ AUTHOR INFORMATION

### Corresponding Authors

Gilles Lemerrier – University of Reims Champagne Ardenne, ICMR, UMR CNRS, F 51687 Reims Cedex 2, France; Email: [gilles.lemerrier@univ-reims.fr](mailto:gilles.lemerrier@univ-reims.fr)

Phong D. Tran – University of Science and Technology of Hanoi, Vietnam Academy of Science and Technology, 100000 Hanoi, Vietnam; [orcid.org/0000-0002-9561-6881](https://orcid.org/0000-0002-9561-6881); Email: [tran.dinh.phong@usth.edu.vn](mailto:tran.dinh.phong@usth.edu.vn)

Catherine Amiens – CNRS, LCC (Laboratoire de Chimie de Coordination), F 31077 Toulouse Cedex 4, France; Université de Toulouse, UPS, INPT, F 31077 Toulouse Cedex 4, France; Email: [catherine.amiens@lcc-toulouse.fr](mailto:catherine.amiens@lcc-toulouse.fr)

### Authors

Quyen T. Nguyen – CNRS, LCC (Laboratoire de Chimie de Coordination), F 31077 Toulouse Cedex 4, France; Université de Toulouse, UPS, INPT, F 31077 Toulouse Cedex 4, France; University of Science and Technology of Hanoi, Vietnam Academy of Science and Technology, 100000 Hanoi, Vietnam

Elodie Rousset – University of Reims Champagne Ardenne, ICMR, UMR CNRS, F 51687 Reims Cedex 2, France

Van T. H. Nguyen – University of Science and Technology of Hanoi, Vietnam Academy of Science and Technology, 100000 Hanoi, Vietnam

Vincent Colliere – CNRS, LCC (Laboratoire de Chimie de Coordination), F 31077 Toulouse Cedex 4, France; Université de Toulouse, UPS, INPT, F 31077 Toulouse Cedex 4, France

Pierre Lecante – CEMES CNRS, Université de Toulouse, F 31055 Toulouse, France

Wantana Klysubun – Synchrotron Light Research Institute, 30000 Nakhon Ratchasima, Thailand

Karine Philippot – CNRS, LCC (Laboratoire de Chimie de Coordination), F 31077 Toulouse Cedex 4, France; Université de Toulouse, UPS, INPT, F 31077 Toulouse Cedex 4, France; [orcid.org/0000-0002-8965-825X](https://orcid.org/0000-0002-8965-825X)

Jérôme Esvan – CIRIMAT, Université de Toulouse, CNRS INPT UPS, 31030 Toulouse, France

Marc Respaud – Université de Toulouse, UPS, INPT, F 31077 Toulouse Cedex 4, France; LPCNO, INSA, 31077 Toulouse Cedex 4, France

Complete contact information is available at: <https://pubs.acs.org/doi/10.1021/acsami.1c15051>

### Author Contributions

Conceptualization—P.D.T., G.L., and C.A.; data curation—E.R., P.L., J.E., V.C., and M.R.; formal analysis—Q.T.N.; funding acquisition—P.D.T., C.A., and K.P.; investigation—

E.R., Q.T.N., W.K., P.L., and V.T.H.N.; methodology—C.A. and P.D.T.; project administration—Q.T.N., C.A., and P.D.T.; supervision—P.D.T., G.L., E.R., C.A., and K.P.; validation—Q.T.N.; visualization—Q.T.N., E.R., M.R., J.E., and P.L.; writing—original draft—Q.T.N., C.A., E.R., P.L., and J.E.; writing—review and editing—Q.T.N., E.R., V.C., P.L., W.K., J.E., M.R., K.P., G.L., P.D.T., and C.A.

## Notes

The authors declare no competing financial interest.

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